publisher, city and/or country where published.

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
(Not for submission under 37 CFR 1.99)
(Not for Submission under or of it may

Application Number		
Filing Date		2006-08-11
First Named Inventor Dimit		rios Pantelidis
Art Unit		
Examiner Name		
Attorney Docket Numb	er	53233-00021 NAT

					_						
U.S.PATENTS Remove											
Examiner Initial*	Cite No	Patent Number	Kind Code ¹	Issue D	Date	Name of Pate of cited Docu	Releva	ges,Columns,Lines where levant Passages or Relevant jures Appear			
	1										
If you wish to add additional U.S. Patent citation information please click the Add button. Add											
U.S.PATENT APPLICATION PUBLICATIONS Remove											
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publica Date	tion	of cited Document			ges,Columns,Lines where levant Passages or Relevant ures Appear		
	1										
If you wish	h to a	dd additional U.S. Publis	hed Ap	plication	citatio	n information p	lease click the Ad	d button	Add		
FOREIGN PATENT DOCUMENTS Remove											
Examiner Initial*	Cite No	Foreign Document Number ³ Country Code ² i Code ⁴ Date Name of Patentee or Applicant of cited Document		or	vhere Rel	or Relevant	T5				
	1	2005/000740	wo		A2	2005-01-06	K.U. Leuven Reser Development	arch &			
If you wish to add additional Foreign Patent Document citation information please click the Add button Add											
-			NON	-PATEN	NT LITE	RATURE DO	CUMENTS	_	Remove		_
Examiner	Examiner Cite Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item								T5		

	Application Number		
NEODMATION DIOCE COURT	Filing Date		2006-08-11
INFORMATION DISCLOSURE STATEMENT BY APPLICANT	First Named Inventor	med Inventor Dimitrios Pantelidis	
Not for submission under 37 CFR 1.99)	Art Unit		
,	Examiner Name		
	Attorney Docket Numb	er	53233-00021 NAT

	1			
If you wisl	h to a	d additional non-patent literature document citation information please click the Add button	Add	

EXAMINER SIGNATURE

Examiner Signature Date Considered

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

See Kind Codes of USPTO Patent Documents at invent_ISPTO_GDI/cr MEPF 9619A. I Enter office that issued the document, by the involved owlf WIPO Standard ST3.) For Linguistee patent document, the origination of the year of the region of the Emperor many precedure the serial number of the patent document, and occurrent by the appropriate symbols as endicated on the document under WIPO Standard ST1.6 If possible, "Applicant is to place a check mark here it regions are required to the patent document."

Application Number Fing Date 2006-08-11 Find Date 2006-08-11 Find Manage Inventor Damtion Parteids At Unit Examiner Name Inventor Damtion Parteids At Unit Examiner Name Attorney Docket Number S2233-00021 NAT

CERTIFICATION STATEMENT

Please see 37	CFR 1.97	and 1.98 to	make the appro	priate selection(s):
---------------	----------	-------------	----------------	----------------------

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 3.7 CFR 1.97(eVI.)

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no tem of information contained in the information disclosure statement was known to any individual designated in 37 CFR 156(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(s)(c)

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- 7 None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

_			
Signature	/C. Rachal Winger/	Date (YYYY-MM-DD)	2006-08-11
Name/Print	C. Rachal Winger	Registration Number	55815

This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file fand by the USPTO to process) an application. Confidentially is governed by \$3 U.S. C.12 and 37 CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patient and Tradenian's (Office, U.S. Operatment of Commence, P. 0. Bast 1436, Alexandria, V.3211.1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.32211.1450.

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested to the patient process and/or examine your submission, which may related that the patient process and/or examine your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pursuant to 5 U.S.C. 552a(m.).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uting an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2014 and 2016. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant for the control of t
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via flori of mapplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patient.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.